

OIPE

RAW SEQUENCE LISTING

DATE: 01/26/2002

PATENT APPLICATION: US/09/730,174

TIME: 13:53:51

Input Set : N:\EBONY'S\ES.txt

Output Set: N:\CRF3\01262002\I730174.raw

Does Not Comply
Corrected Diskette Needs

3 <110> APPLICANT: R.J. ZAHRADNIK

W--> 4 <120> TITLE OF INVENTION: ANTIBODIES AND PEPTIDE ANTIGENS FOR PRODUCING ANTIBODIES HAVING A

W--> 5 SELECTIVE BINDING SPECIFICITY TO BIOACTIVE INTACT PARATHYROID HORMONE (PTH) 1 84

W--> 6 <130> FILE REFERENCE: IMUNE-001A

W--> 7 <140> CURRENT APPLICATION NUMBER: 09/730,174

C--> 8 <141> CURRENT FILING DATE: 2002-01-09

E--> 8 <160> NUMBER OF SEQ ID: 12

following pages

ERRORED SEQUENCES

- 10 <210> SEQ ID NO: 1
- 11 <211> LENGTH: 11
- 12 <212> TYPE: PRT
- 13 <213> ORGANISM: ARTIFICIAL SEQUENCE
- W--> 14 <220> FEATURE: INDUCES FORMATION OF ANTIBODIES AND ISOLATES SAID ANTIBODIES
- W--> 14 <220> FEATURE: INDUCES FORMATION OF ANTIBODIES AND ISOLATES SAID ANTIBODIES
 - 15 <223> OTHER INFORMATION:
- W--> 16 <400> SEQUENCE: 001; VAL SER GLU ILE GLN LEU MET HIS ASN LEU GLY
- E--> 16 001; VAL SER GLU ILE GLN LEU MET HIS ASN LEU GLY
 - 18 <210> SEQ ID NO: 2
 - 19 <211> LENGTH: 11
 - 20 <212> TYPE: PRT
 - 21 <213> ORGANISM: ARTIFICIAL SEQUENCE
- W--> 22 <220> FEATURE: INDUCES FORMATION OF ANTIBODIES AND ISOLATES SAID ANTIBODIES
- W--> 22 <220> FEATURE: INDUCES FORMATION OF ANTIBODIES AND ISOLATES SAID ANTIBODIES
 - 23 <223> OTHER INFORMATION:
- W--> 24 <400> SEQUENCE: 002; VAL SER GLU ILE GLN PHE MET HIS ASN LEU GLY
- E--> 24 002; VAL SER GLU ILE GLN PHE MET HIS ASN LEU GLY
 - 26 < 210 > SEQ ID NO: 3
 - 27 <2115 LENGTH: 12
 - 28 <212> TYPE: PRT
 - 29 <213> ORGANISM: ARTIFICIAL SEQUENCE
- w--> 30 <220> FEATURE: INDUCES FORMATION OF ANTIBODIES AND ISOLATES SAID ANTIBODIES
- W--> 30 <220> FEATURE: INDUCES FORMATION OF ANTIBODIES AND ISOLATES SAID ANTIBODIES
 - 31 <223> OTHER INFORMATION:
- W--> 32 <400> SEQUENCE: 003; SER VAL SER GLU ILE GLN LEU MET HIS ASN LEU GLY
- E--> 32 003; SER VAL SER GLU ILE GLN LEU MET HIS ASN LEU GLY
 - 34 <210> SEQ ID NO: 4
 - 35 <211> LENGTH: 12
 - 36 <:212> TYPE: PRT
 - 37 <213> ORGANISM: ARTIFICIAL SEQUENCE
- W--> 38 <220> FEATURE: INDUCES FORMATION OF ANTIBODIES AND ISOLATES SAID ANTIBODIES

SEQUENCE-LISTING-

-9EAABIM	OB-DIOTINO -	
<110>	R.J. ZAHRADNIK	
<120>	ANTIBODIES AND PEPTIDE ANTIGENS FOR PRODUCING ANTIBODIES HAVING A	. 1 04
	IVE BINDING SPECIFICITY TO BIOACTIVE INTACT PARATHYROID HORMONE (PTH	•
<130>	IMUNE-001A	rd response
<140>	1MUNE-001A 09/730, 174 12 12 14/7 C-ensert this numeric electific a	
<160>	12	
<210>	1	
<211>	11	
<212>	PRT	
<213>	ARTIFICIAL SEQUENCE	, ,
<220>	INDUCES FORMATION OF ANTIBODIES AND ISOLATES SAID ANTIBODIES 1920	re Anteretur
<223> <i>&</i>	-1. It wast hard return much a mere winder to last to last	
<400>	001 July insert hard return more amuse ands to line below you	10-62237 line
	1 5. 10	Do NOT west
<210>	2 number the arrest and white every 5 amend 11 PRT (PC NUT we TAB codes)	DO NOT LASTA
<211>	11	(1 . 44 . 44 . 244
<212>	PRT (PO NUT Use TAB Codes)	and respective
<213>	ADMITTATAL GEOMENCE	
	ARTIFICIAL SEQUENCE INDUCES FORMATION OF ANTIBODIES AND ISOLATES SAID ANTIBODIES	Lo (220%.
<220>		
<223>	hard return	22257 Min
<400>	0026 VAL SER GLU ILE GLN PHE MET HIS ASN LEU GLY	120 0 1
	$1 \sim 1$	header
<210>	3	27.14
<211>	12	only.
<212>	PRT	V
<213>	ARTIFICIAL SEQUENCE	1121
<220>	INDUCES FORMATION OF ANTIBODIES AND ISOLATES SAID ANTIBODIES	0020
<223>		Lunne-case
<400>	-CR23() & SER VAL SER GLU ILE GLN LEU MET HIS ASN LEU GLY	letter for
	1 5 10	0. 1101/0
<210>	4	secon jui
<211>	12	71
<212>	PRT	the initial
<213>	ARTIFICIAL SEQUENCE	
<220>	INDUCES FORMATION OF ANTIBODIES AND ISOLATES SAID ANTIBODIES.	letter of
<223>	is hard return)
< 400>	467 SER VAL SER GLU ILE GLM PHE MFT HIS ASN LEU GLY	amero and
	1 5 /0	
<210>	5	0. V.1
<211>	12	eg. val
<212>	PRT	
<213>	ARTIFICIAL SEQUENCE	Σεr
< 32.00	INDUCES FORMATION OF ANTIBODIES AND ISOLATES SAID ANTIBODIES	eg. Val Ser Glu Ile Cln
< 7.7 % x	el hald eleven	<u>U</u> /U
< 4000>	561 ALA VAL SER GLU ILE GLN LEU MET HIS ASN LEU GLY	71,
	1 5 /0	<u></u>
<2105	6	().
<211>	12	UIN
(,,	PRT	=
<.13.	ARTIFICIAL SEQUENCE	NOT VAL
<:220>	INDUCES FORMATION OF ANTIBODIES AND ISOLATES SAID ANTIBODIES	NOTVAL
<223>	hard return	(= 0
<400>	6; 4 ALA VAL SER GLU ILE GLN PHE MET HIS ASN LEU GLY	SER
44002		6111
	A Prose	GLU
	, the same	L.
	and a delication of the second	tracted
	(samue of years) ones) sequered listing (Charles Car /
	(sample o) global orrers) Please consult sample for valid form	at
	The state for the	• /

<110>	Smith, John: Smithgene Inc.						
<120>	Example of a Sequence Listing						
<130>	01-00001						
<140>	PCT/EP98/00001						
<1(1)	1998-12-31						
<150> <151>	US 08/999,999 1997-10-15						
<160>	*						
<170>	PatentIn version 2.0	•					
<210>	1						
<211>	389						
<212>	DNA 🤔						
<213>	Paramecium sp.						
<220>							
< 221>	CDS						
<222>	(279)(389)						
<300>							
<301>	Doe, Richard						
<302>	Isolation and Characterization of a Gene Encoding a						
	Protease from Paramecium sp.						
< 303>	Journal of Genes						
<304>	1						
<305>	4						
< 306 >	1 - 7						
< 307>	1988-06-31						
< 308>	123456						
< 309>	1988-06-31						
< 400 >							
agetglagic	allocigigi colollolot olgggolici caccolgola alcagatoto	60					
	and the state of t						
agggagagtg	E CONTRACTOR DE LA CONT	20					
rgargrggca	attgotggca gtgndadayy სამანასაფიი აჭვინნაფიი ნფფინნ cc gc l	80					
cgcggcgcgg	eggecectet egegeteete tegegeetet etetegetet eeletegete 2	4 0					

Please consult

I

Appendix 3, page 2

ggac	ctga 	tt a	iggtga	gcag	9899	39999	Ca	gttago		atg Met 1	gtt Val	tca Ser	atg Met	ttc Phe S	agc Ser	296
ttg Leu	tct Ser	t t c Phe		tgg Trp	cct Pro	gga Gly	ttt Phe	tgt Cys 15	ttg Leu	ttt Phe	gtt Val	tgt Cys	ttg Leu 20	ttc Phe	caa Cln	344
tgt Cys	ccc Pro	aaa Lys 25	otc -Val	ctc Lcu	ccc Pro	tgt Cys	cac His	t ca Ser	tca Ser	ctg -Lcu	cag Gln	ccg Pro 35		ctt L bi u		389
<210: <211: <212: <213:	> >			ium s	ρ.			<i>:</i>	15	· · ·	7			yc:		
<4002 Het 1	Val	2 Ser	Het	Phe 5	Sër	Leu	Ser	Phe	Lys 10	Trp	Pro	Cly	Phg	Cys 15	Leu	
Phe	Val	Cys	Leu 20	Phe	Cln	Cys	Pro	Lys 25	Va l	Leu	Pro	Cys	His 30	Ser	Ser .	
Leu	Gln	Pro 35	ksn	L q u												
<210><211><211><212><213>		} 11 PR ^r		al Sc	quenc	c								•		
<220> <223>		De Lis	signec nker l	l pepu	ide b n the	ased c	on si:	ze and beta	pola chair	rity is of	to ac Prote	tas d in XY	a Z .			
<400> Net l	Val) Asn	Leu	Glu S	Pro	Met	His	Thr (Clu 10	lle						
<210><400>		4														

(Annex VIII follows)

E

table. The numeric identifier shall be used only in the 'Sequence Listing." The order and presentation of the items of information in the "Sequence Listing" shall conform to the arrangement given below. Each item of information shall begin on a new line and shall begin with the numeric identifier enclosed in angle brackets as shown. The submission of those items of information designated with an "M" is mandatory. The submission of those items of information designated with an "O" is optional. Numeric identifiers <110> through <170> shall only be set forth at the beginning of the "Sequence Listing." The following table illustrates the numeric identifiers.

Numeric Identifier	Definition	Comments and Format	Mandatory (M) or Optional (O)			
<110>	Applicant	Preferably max. of 10 names; one name per line; preferable format: Surname, Other: Names and/or Initials	M W			
<120>	Title of Invention		M			
<130>	File Reference	Personal file reference	M, when filed prior to assignment of appl. number			
<140>	Current Applica- tion Number	Specify as: US 07/999,999 or PCT/US96/99999	M, if available			
< 1 4 1 >	Current Filing Date	Specify as: yyyy-mm-dd	M, if available			
<150>	Prior Application Number	Specify as: US 07/999,999 or PCT/US96/99999	M, if applicable include priority documents under 35 USC 119 and 120			
<151>	Prior Application Filing Date	Specify as: yyyy-mm-dd	M, if applicable			
<160>	Number of SEQ ID	Count includes total number of SEQ ID NOs	М			
<170>	Software	Name of software used to create the Sequence Listing	0			
<210>	SEQ ID NO: # .	Response shall be an integer representing the SEQ ID NO shown	м			
<211>	Length	Respond with an integer expressing the number of bases or amino acid residues	м			

Whether presented sequence moleculc is DNA, RNA, or PRT (protein). If a nucleotide sequence contains both DNA and RNA fragments, the type shall be "DNA." In addition, the combined DNA/ RNV wolcculc shall be further described in

<213>

Organism

Scientific name, i.e. Genus/species, Unknown or Artificial Sequence. In addition, the "Unknown" or "Artificial Sequence" organisms shall be further described in the <220> to <223> feature section.

the <220> to

<223> feature section.

<220>

Feature

Leave blank after <220>. <221-223> provide for a description of points of biological significance in the sequence.

M, under the following conditions: if "n,"
"Xaa," or a modified or unusual L-amino acid or modified base was used in a sequence; if ORGANISM is "Artificial Sequence" or "Unknown"; if molecule is combined DNA/RNA.

نتو

<221>

Name/Key

Provide appropriate identifier for feature, preferably from wiPO Standard ST.25 (1998), Appendix 2, Tables 5 and 6

M, under the following conditions: = if "n," "Xaa," or a modified or unusual L-amino acid or modified base was used in a sequence

<222>

Location

Specify location within sequence; where appropriate state number of first and last bases/amino acids

M, under the following conditions:
if "n," "Xaa," or
a modified or un
usual L-amino
acid or modified

in feature

ise was used in a sequence

<223>	Other Information	Other relevant information; four lines maximum	M, under the following conditions: if "n," "Xaa," or a modified or unusual L-amino acid or modified base was used in a sequence; if ORGANISM is "Artificial Sequence" or "Unknown"; if and acid molecule is combined DNA/RNA.
<300>	Publication Information	Leave blank after <300> /*	0
<301>	Authors	Preferably max of ten named authors of publication; specify one name per line; preferable format: Surname, Other Names and/or Initials	• • • •
<302>	Title		0
<303>	Journal		0
< 304 >	Volume		0
<305>	Issue		0
< 306>	Pages		0 ·
<307>	Date	Journal date on which data published; specify as yyyy-mm- dd, MMM-yyyy or Season-yyyy	O
<300>	Database Accession Number	Accession number assigned by data- base including database name	O E
<309;	Database Entry Date	Date of entry in database: specity as yyyy:mm-dd or MMM-yyyy	0
<310>	Patent Document Number	Document number; for patent-type citations only. Specify as, for example, US 07/999,999	0

* <311*.	Patent Firing Date	Document filing date, for patent-type citations only; specify as yyyy-mm-dd		
<312>	Publication Date	Document publication date, for patent-type citations only; specify as yyyy-mm-dd-	o .	
<313>	Relevant Residues	FROM (position) TO (position)	0	•
<400>	Sequence	follow the numeric identifier and should appear on the line pre-		æ å æ, æ

- 5. Section 1.024 is revised to read as follows:
- 1.824 Form and format for nucleotide and/or amino acid sequence submissions in computer readable form.
- (a) The computer readable form required by 1.821(e) shall meet the following specifications:
- (1) The computer readable form shall contain a single "Sequence Listing" as either a diskette, series of diskettes, or other permissible media outlined in paragraph (c) of this section.
- (2) The "Sequence Listing" in paragraph (a) (1) of this section shall be submitted in American Standard Code for Information Interchange (ASCII) text. No other formats shall be allowed.
- (3) The computer readable form may be created by any means, such as word processors, nucleotide/amino acid sequence editors or other custom computer programs; however, it shall conform to all specifications detailed in this section.
- (4) File compression is acceptable when using diskette media, so long as the compressed file is in a self-extracting format that will decompress on one of the systems described in paragraph (b) of this section.
- (5) Page numbering shall not appear within the computer readable form version of the "Sequence Listing" (ile.
- (6) All computer readable forms shall have a label permanently affixed thereto on which has been hand-printed or typed: the name of the applicant, the title of the invention, the date on which the data were recorded on the computer readable form, the operating system used, a reference number, and an application serial number and filing date, if known.
- (b) Computer readable form submissions must meet these format requirements:
- (I) Computer: IBM PC/XT/AT, or compatibles, or Apple Macintosh;
- (2) Operating System: MS-DOS, Unix or Macintosh;

1.1

0590



...



The Biotechnology Systems Branch of the Scientific and Technical Information Center (STIC) detected errors when processing the following computer readable form:

09/730,174
O'IPE,
1/26/2002

THE ATTACHED PRINTOUT EXPLAINS DETECTED ERRORS.
PLEASE FORWARD THIS INFORMATION TO THE APPLICANT BY EITHER:

- 1) INCLUDING A COPY OF THIS PRINTOUT IN YOUR NEXT COMMUNICATION TO THE APPLICANT, WITH A NOTICE TO COMPLY or,
- 2) TELEPHONING APPLICANT AND FAXING A COPY OF THIS PRINTOUT, WITH A NOTICE TO COMPLY

FOR CRF SUBMISSION QUESTIONS, PLEASE CONTACT MARK SPENCER, 703-308-4212.

FOR SEQUENCE RULES INTERPRETATION, PLEASE CONTACT ROBERT WAX, 703-308-4216. PATENTIN 2.1 e-mail help: patin21help@uspto.gov or phone 703-306-4119 (R. Wax) PATENTIN 3.0 e-mail help: patin3help@uspto.gov or phone 703-306-4119 (R. Wax)

TO **REDUCE** ERRORED SEQUENCE LISTINGS, **PLEASE** USE THE <u>CHECKER</u> <u>VERSION 3.1 PROGRAM</u>, ACCESSIBLE THROUGH THE U.S. PATENT AND TRADEMARK OFFICE WEBSITE. SEE BELOW FOR ADDRESS:

http://www.uspto.gov/web/offices/pac/checker

Applicants submitting genetic sequence information electronically on diskette or CD-Rom should be aware that there i a possibility that the disk/CD-Rom may have been affected by treatment given to all incoming mail. Please consider using alternate methods of submission for the disk/CD-Rom or replacement disk/CD-Rom. Any reply including a sequence listing in electronic form should NOT be sent to the 20231 zip code address for the United States Patent and Trademark Office, and instead should be sent via the following to the indicated addresses:

- 1. EFS-Bio (http://www.uspto.gov/ebc/efs/downloads/documents.htm, EFS Submission User Manual ePAVE)
- 2. U.S. Postal Service: U.S. Patent and Trademark Office, Box Sequence, P.O. Box 2327, Arlington, VA 22202
- 3. Hand Carry directly to:
 - U.S. Patent and Trademark Office, Technology Center 1600, Reception Area, 7th Floor, Examiner Name, Sequence Information, Crystal Mall One, 1911 South Clark Street, Arlington, VA 22202
 - U.S. Patent and Trademark Office, Box Sequence, Customer Window, Lobby, Room 1B03, Crystal Plaza Two, 2011 South Clark Place, Arlington, VA 22202
- 4. Federal Express, United Parcel Service, or other delivery service to: U.S. Patent and Trademark Office, Box Sequence, Room 1B03-Mailroom, Crystal Plaza Two, 2011 South Clark Place, Arlington, VA 22202

Revised 01/29/2002